

Message

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**From:** Miller, David [/O=EXCHANGELABS/OU=EXCHANGE ADMINISTRATIVE GROUP (FYDIBOHF23SPDLT)/CN=RECIPIENTS/CN=FA0582F5BA6540C687844F9289A4F74F-DAVID J. MILLER]  
**Sent:** 12/14/2020 4:56:26 PM  
**To:** Doherty, Michael [Doherty.Michael@epa.gov]  
**CC:** Niman, Aaron (niman.aaron@epa.gov) [niman.aaron@epa.gov]  
**Subject:** FW: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

And this...

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**From:** Miller, David  
**Sent:** Thursday, January 16, 2020 10:05 AM  
**To:** Doherty, Michael <Doherty.Michael@epa.gov>  
**Cc:** Niman, Aaron (niman.aaron@epa.gov) <niman.aaron@epa.gov>  
**Subject:** FW: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

I knew that is what you meant ; - )

I was curious as to what your correction to Don's interpretation would. I can't imagine the CCPR Secretariat wanting to touch this issue with a 10 foot pole. The folks that think they would or might are operating in a fantasy land. And having us operate in a fantasy land where nobody officially makes a request so nobody has to officially deny a request helps Dr. Hodges and he is happy we are there and to not disavow us of our notions.

David.

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**From:** Doherty, Michael  
**Sent:** Thursday, January 16, 2020 10:00 AM  
**To:** Sadowsky, Don <Sadowsky.Don@epa.gov>; Miller, David <Miller.DavidJ@epa.gov>; Ingram, Earl <Ingram.Earl@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>  
**Cc:** Furlow, Calvin <Furlow.Calvin@epa.gov>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Hi Don,

## Ex. 5 Attorney Client (AC)

Mike

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**From:** Sadowsky, Don <Sadowsky.Don@epa.gov>  
**Sent:** Thursday, January 16, 2020 9:56 AM  
**To:** Doherty, Michael <Doherty.Michael@epa.gov>; Miller, David <Miller.DavidJ@epa.gov>; Ingram, Earl <Ingram.Earl@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>  
**Cc:** Furlow, Calvin <Furlow.Calvin@epa.gov>  
**Subject:** Re: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

## Ex. 5 Attorney Client (AC)

# Ex. 5 Attorney Client (AC)

## Ex. 5 Attorney Client (AC)

Donald A. Sadowsky  
Pesticides and Toxic Substances Law Office  
Office of General Counsel  
U.S. Environmental Protection Agency  
1200 Pennsylvania Avenue, N.W.  
Washington, DC 20460  
(202) 564-5638

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**From:** Doherty, Michael <[Doherty.Michael@epa.gov](mailto:Doherty.Michael@epa.gov)>  
**Sent:** Thursday, January 16, 2020 9:45 AM  
**To:** Miller, David <[Miller.DavidJ@epa.gov](mailto:Miller.DavidJ@epa.gov)>; Sadowsky, Don <[Sadowsky.Don@epa.gov](mailto:Sadowsky.Don@epa.gov)>; Ingram, Earl <[Ingram.Earl@epa.gov](mailto:Ingram.Earl@epa.gov)>; Niman, Aaron <[niman.aaron@epa.gov](mailto:niman.aaron@epa.gov)>; Johnson, Marion <[Johnson.Marion@epa.gov](mailto:Johnson.Marion@epa.gov)>  
**Cc:** Furlow, Calvin <[Furlow.Calvin@epa.gov](mailto:Furlow.Calvin@epa.gov)>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

## Ex. 5 Attorney Client (AC)

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**From:** Miller, David <[Miller.DavidJ@epa.gov](mailto:Miller.DavidJ@epa.gov)>  
**Sent:** Wednesday, January 15, 2020 2:31 PM  
**To:** Sadowsky, Don <[Sadowsky.Don@epa.gov](mailto:Sadowsky.Don@epa.gov)>; Ingram, Earl <[Ingram.Earl@epa.gov](mailto:Ingram.Earl@epa.gov)>; Niman, Aaron <[niman.aaron@epa.gov](mailto:niman.aaron@epa.gov)>; Johnson, Marion <[Johnson.Marion@epa.gov](mailto:Johnson.Marion@epa.gov)>  
**Cc:** Furlow, Calvin <[Furlow.Calvin@epa.gov](mailto:Furlow.Calvin@epa.gov)>; Doherty, Michael <[Doherty.Michael@epa.gov](mailto:Doherty.Michael@epa.gov)>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

# Ex. 5 Attorney Client (AC)

David

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**From:** Sadowsky, Don  
**Sent:** Wednesday, January 15, 2020 2:29 PM  
**To:** Miller, David <[Miller.DavidJ@epa.gov](mailto:Miller.DavidJ@epa.gov)>; Ingram, Earl <[Ingram.Earl@epa.gov](mailto:Ingram.Earl@epa.gov)>; Niman, Aaron <[niman.aaron@epa.gov](mailto:niman.aaron@epa.gov)>; Johnson, Marion <[Johnson.Marion@epa.gov](mailto:Johnson.Marion@epa.gov)>  
**Cc:** Furlow, Calvin <[Furlow.Calvin@epa.gov](mailto:Furlow.Calvin@epa.gov)>; Doherty, Michael <[Doherty.Michael@epa.gov](mailto:Doherty.Michael@epa.gov)>  
**Subject:** Re: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

## Ex. 5 Attorney Client (AC)

Donald A. Sadowsky  
Pesticides and Toxic Substances Law Office  
Office of General Counsel

U.S. Environmental Protection Agency  
1200 Pennsylvania Avenue, N.W.  
Washington, DC 20460  
(202) 564-5638

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**From:** Miller, David <Miller.DavidJ@epa.gov>  
**Sent:** Wednesday, January 15, 2020 2:26 PM  
**To:** Ingram, Earl <Ingram.Earl@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>  
**Cc:** Furlow, Calvin <Furlow.Calvin@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Sadowsky, Don <Sadowsky.Don@epa.gov>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Thanks, Earl. Not necessarily unexpected.

I will forward to you all as an FYI what I sent to all or most of you back in November clarify some of the implications that were portrayed in Dr. Hodges email note to the Codex Secretariat. Specifically, Dr. Hodges seemed to imply that it was EPA that was the hold-up in this process of getting the studies released.

At some point we will want to discuss where we want to be on this when Aaron and I attend the CCPR meeting in April. As it stands now, CCPR is being led to believe by Dr. Hodges's earlier email note that EPA will be forthcoming with these required studies in sufficient time for aldicarb to undergo its (now rescheduled) Codex periodic review in September 2021.

David.

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**From:** Ingram, Earl  
**Sent:** Wednesday, January 15, 2020 2:07 PM  
**To:** Niman, Aaron <niman.aaron@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>  
**Cc:** Furlow, Calvin <Furlow.Calvin@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Sadowsky, Don <Sadowsky.Don@epa.gov>; Miller, David <Miller.DavidJ@epa.gov>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Aaron,

They have not submitted a FOIA request.

Earl Ingram, Chief  
Public Information & Records Integrity Branch  
Information Technology & Resources Management Division  
Office of Pesticide Programs  
U.S. Environmental Protection Agency  
(703) 305-5456

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**From:** Niman, Aaron <niman.aaron@epa.gov>  
**Sent:** Wednesday, January 15, 2020 1:31 PM  
**To:** Ingram, Earl <Ingram.Earl@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>  
**Cc:** Furlow, Calvin <Furlow.Calvin@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Sadowsky, Don <Sadowsky.Don@epa.gov>; Miller, David <Miller.DavidJ@epa.gov>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Earl and Marion,

David asked me to follow-up on the status of aldicarb to determine if AgLogic has submitted a FOIA request to obtain aldicarb data owned by Bayer. If you recall, aldicarb is subject to periodic review by the Joint FAO/WHO Meeting on Pesticide Residues (JMPR) and is seeking the data to support JMPR's periodic review process.

Have you received any new information or requested from Larry Hodges of AgLogic?

Thanks,

-Aaron

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**From:** Ingram, Earl <[Ingram.Earl@epa.gov](mailto:Ingram.Earl@epa.gov)>  
**Sent:** Monday, November 4, 2019 1:56 PM  
**To:** Miller, David <[Miller.DavidJ@epa.gov](mailto:Miller.DavidJ@epa.gov)>  
**Cc:** Furlow, Calvin <[Furlow.Calvin@epa.gov](mailto:Furlow.Calvin@epa.gov)>; Niman, Aaron <[niman.aaron@epa.gov](mailto:niman.aaron@epa.gov)>; Doherty, Michael <[Doherty.Michael@epa.gov](mailto:Doherty.Michael@epa.gov)>; Sadowsky, Don <[Sadowsky.Don@epa.gov](mailto:Sadowsky.Don@epa.gov)>; Johnson, Marion <[Johnson.Marion@epa.gov](mailto:Johnson.Marion@epa.gov)>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

We have not received a FOIA request from Mr. Hodges.

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**From:** Miller, David <[Miller.DavidJ@epa.gov](mailto:Miller.DavidJ@epa.gov)>  
**Sent:** Monday, November 04, 2019 1:27 PM  
**To:** Ingram, Earl <[Ingram.Earl@epa.gov](mailto:Ingram.Earl@epa.gov)>  
**Cc:** Furlow, Calvin <[Furlow.Calvin@epa.gov](mailto:Furlow.Calvin@epa.gov)>; Niman, Aaron <[niman.aaron@epa.gov](mailto:niman.aaron@epa.gov)>; Doherty, Michael <[Doherty.Michael@epa.gov](mailto:Doherty.Michael@epa.gov)>; Sadowsky, Don <[Sadowsky.Don@epa.gov](mailto:Sadowsky.Don@epa.gov)>; Johnson, Marion <[Johnson.Marion@epa.gov](mailto:Johnson.Marion@epa.gov)>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Ok. Thank you.

And can I or can I not assume that ITRMD has such formal FOIA request in hand from Dr. Hodges as per Don S's Item (1) below from September?

David

---

**From:** Ingram, Earl <[Ingram.Earl@epa.gov](mailto:Ingram.Earl@epa.gov)>  
**Sent:** Monday, November 04, 2019 1:14 PM  
**To:** Miller, David <[Miller.DavidJ@epa.gov](mailto:Miller.DavidJ@epa.gov)>  
**Cc:** Furlow, Calvin <[Furlow.Calvin@epa.gov](mailto:Furlow.Calvin@epa.gov)>; Niman, Aaron <[niman.aaron@epa.gov](mailto:niman.aaron@epa.gov)>; Doherty, Michael <[Doherty.Michael@epa.gov](mailto:Doherty.Michael@epa.gov)>; Sadowsky, Don <[Sadowsky.Don@epa.gov](mailto:Sadowsky.Don@epa.gov)>; Johnson, Marion <[Johnson.Marion@epa.gov](mailto:Johnson.Marion@epa.gov)>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Yes.

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**From:** Miller, David <[Miller.DavidJ@epa.gov](mailto:Miller.DavidJ@epa.gov)>  
**Sent:** Monday, November 04, 2019 12:54 PM  
**To:** Ingram, Earl <[Ingram.Earl@epa.gov](mailto:Ingram.Earl@epa.gov)>  
**Cc:** Furlow, Calvin <[Furlow.Calvin@epa.gov](mailto:Furlow.Calvin@epa.gov)>; Niman, Aaron <[niman.aaron@epa.gov](mailto:niman.aaron@epa.gov)>; Doherty, Michael <[Doherty.Michael@epa.gov](mailto:Doherty.Michael@epa.gov)>; Sadowsky, Don <[Sadowsky.Don@epa.gov](mailto:Sadowsky.Don@epa.gov)>; Johnson, Marion <[Johnson.Marion@epa.gov](mailto:Johnson.Marion@epa.gov)>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

So.... Was the information below suggested by Don Sadowsky actually conveyed to Dr. Hodges by OPP ITRMD?

David

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**From:** Ingram, Earl <Ingram.Earl@epa.gov>  
**Sent:** Monday, November 04, 2019 12:38 PM  
**To:** Miller, David <Miller.DavidJ@epa.gov>  
**Cc:** Furlow, Calvin <Furlow.Calvin@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Sadowsky, Don <Sadowsky.Don@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

You are welcome.

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**From:** Miller, David <Miller.DavidJ@epa.gov>  
**Sent:** Monday, November 04, 2019 11:54 AM  
**To:** Ingram, Earl <Ingram.Earl@epa.gov>  
**Cc:** Furlow, Calvin <Furlow.Calvin@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Sadowsky, Don <Sadowsky.Don@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>  
**Subject:** FW: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Thanks, Earl. This is helpful.

I see that Don suggests in his September email the following:

## Ex. 5 Attorney Client (AC)

Was the above information conveyed to Dr. Hodges? If so: (A) it seems EPA is being fairly clear with Dr. Hodges; and (B) Dr. Hodges is not being clear with Codex.

David.

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**From:** Ingram, Earl <Ingram.Earl@epa.gov>  
**Sent:** Monday, November 04, 2019 11:31 AM  
**To:** Miller, David <Miller.DavidJ@epa.gov>; Furlow, Calvin <Furlow.Calvin@epa.gov>  
**Cc:** Niman, Aaron <niman.aaron@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>; Metzger, Michael <Metzger.Michael@epa.gov>; Sadowsky, Don <Sadowsky.Don@epa.gov>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

David,

## Ex. 5 Attorney Client (AC)

I have copied Don.

Earl

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**From:** Miller, David <Miller.DavidJ@epa.gov>  
**Sent:** Monday, November 04, 2019 11:15 AM  
**To:** Furlow, Calvin <Furlow.Calvin@epa.gov>; Ingram, Earl <Ingram.Earl@epa.gov>  
**Cc:** Niman, Aaron <niman.aaron@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>; Metzger, Michael <Metzger.Michael@epa.gov>  
**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Dear Earl and Calvin,

I read the message below from Mr. Larry Hodges which was sent to you as well as a number of officials from the Codex Secretariat and well as the Australian Chair of the CCPR's electronic Working Group on Priorities. Larry's email below -- whether accurate or not -- seems to suggest that US EPA itself will be providing the studies to CCPR, regardless of what I understand are objections to this from Bayer Crop Science. I am not an attorney and not an expert in 6(a)(2), but it strikes me that EPA's release of this to CCPR would be in violation of the law. If I am wrong in that, I would be happy to be corrected. Have you consulted with OGC on this and have they indicated that EPA's release of this proprietary Bayer data to CCPR is alright? It would be helpful for you to send to me and Aaron Niman any communication you have had with Mr. Hodges indicating that it is US EPA's intention to release proprietary Bayer data to CCPR, data which last I know Bayer had not agreed could be released by the Agency.

With Mr Hodges cc'ing a slew of officials from CCPR with an email that asks the planned upcoming CCPR meeting covering aldicarb to be delayed "to allow enough time for the US EPA to submit the studies to the CCPR", this certainly to me now implies that EPA had agreed to and endorses release of the Bayer data, and puts EPA -- and our US Delegation to CCPR - in a poor light if this data is not later forthcoming. I note that it confuses the CCPR planning and prioritization process and may ultimately displace other legitimate pesticides on the CCPR priority list, thereby potentially delaying their approval.

Mr. Hodges in his email appears to have placed the onus on EPA to provide the Bayer data required by JMPR to Codex. If this is true, it would be useful to hear this from you two directly. If not true, then I believe information that Mr. Hodges provided in his email below needs to be clarified by indicating that EPA has **not** agreed at this time to release the Bayer data to Codex so that any misimpressions can be corrected. What's there now in his Mr. Hodges email certainly gives that impression.

Clarification on this point from the both of you would be appreciated.

Thank you.

Regards,

David.

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David J. Miller CAPT | USPHS  
U.S. Delegate to Codex Committee on Pesticide Residues (CCPR)  
Chief, Chemistry & Exposure Branch  
and Acting Chief, Toxicology & Epidemiology Branch  
Health Effects Division  
Office of Pesticide Programs  
703-305-5352 (voice)  
703-305-5147 (fax)

Visit [www.epa.gov/pesticides](http://www.epa.gov/pesticides)

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**From:** Larry Hodges <larryhodges@meycorp.com>

**Sent:** Monday, November 04, 2019 9:59 AM

**To:** Reichstein, Ian <Ian.Reichstein@agriculture.gov.au>; MADSEN, Soren <madsens@who.int>

**Cc:** VERGER, Philippe <vergerp@who.int>; Yang, YongZhen (AGPM) <YongZhen.Yang@fao.org>; Miller, David <Miller.DavidJ@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Ingram, Earl <IMCEAMAILTO-Ingram+2EEarl+40epa+2Egov@namprd04.prod.outlook.com>; Furlow, Calvin <Furlow.Calvin@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>; Antoine Puech <antoinepuech@meycorp.com>

**Subject:** RE: Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Dear Mr. Reichstein and Mr. Madsen,

As previously explained, AgLogic Chemical (the current and only registrant of aldicarb) is committed to supporting the existing aldicarb MRLs but we do not have access to the aldicarb studies. These studies were submitted to the US EPA by Bayer CropScience and it is unlikely that the US EPA will be able to provide the aldicarb studies to the CCPR by December 1, 2019. As it is extremely important that aldicarb not be removed from the CCPR list of pesticides we propose that the aldicarb reevaluation be rescheduled from 2020 to 2021 to allow enough time for the US EPA to submit the studies to the CCPR.

Please respond to this email and let us know how the CCPR wishes to proceed and what we can do to ensure the necessary aldicarb studies are available for review.

Best Regards,  
Larry Hodges, Ph.D.  
Director of Regulatory Affairs  
AgLogic Chemical LLC

Phone: 919-932-5800

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**From:** Larry Hodges

**Sent:** Thursday, October 10, 2019 1:08 PM

**To:** Reichstein, Ian <Ian.Reichstein@agriculture.gov.au>; MADSEN, Soren <madsens@who.int>

**Cc:** VERGER, Philippe <vergerp@who.int>; YongZhen.Yang@fao.org; Miller, David <Miller.DavidJ@epa.gov>; Niman, Aaron <niman.aaron@epa.gov>; Doherty, Michael <Doherty.Michael@epa.gov>; Ingram, Earl ; Furlow, Calvin <Furlow.Calvin@epa.gov>; Johnson, Marion <Johnson.Marion@epa.gov>; Antoine Puech <antoinepuech@meycorp.com>

**Subject:** Toxicological and Residue Evaluation Aldicarb (117) by CCPR/JMPR

Dear Mr. Reichstein and Mr. Madsen,

I am sending this email to advise you of the current status of the data submission for aldicarb (117) and to ask for guidance on the path forward. The attached two documents, summarized below, provide the background on our attempt to provide the CCPR with the requested aldicarb studies.

On October 24, 2018 AgLogic Chemical LLC was advised by Mr. Ian Reichstein that aldicarb was reinstated as a supported compound awaiting periodic review and would be placed as a confirmed listing for the 2020 Schedule.

We notified the CCPR that AgLogic is a generic registrant and, as allowed by US law, our aldicarb registration is supported by toxicology and residue data that were developed and submitted to the US Environmental Protection Agency (EPA) by aldicarb's previous registrant, Bayer CropScience. We are not allowed to have copies of the actual studies that support registration and, therefore, despite our best intentions, we are not able to submit these studies to the JMPR for review.

We suggested that since the EPA already has the supporting studies, the aldicarb reviews could be written by the EPA's residue and toxicology experts that participate in the JMPR or, alternatively, the EPA could provide the required studies to the JMPR for review by someone not from the EPA.

On May 22, 2019 AgLogic Chemical was notified by Mr. Soren Madsen that the aldicarb studies should be submitted to the JMPR for review in order to avoid possible (or perceived) conflicts of interest. We were told that the JMPR secretariat seeks to assign monographers and reviewers that have not been directly involved in recent national evaluations of the assigned compound.

We immediately contacted the EPA and asked if they could submit the aldicarb studies directly to the JMPR and EPA said that it was most appropriate to work through the Freedom of Information Act (FOIA). That determination was made based upon the requirements set forth under Section 10(g) of FIFRA, which, in part, restricts the availability of how studies/data submitted to the EPA can be released to a specific requestor. At this point the EPA Office of General Counsel contacted Bayer CropScience to get their consent to release the aldicarb studies to the JMPR.

On July 29, 2019 AgLogic Chemical was notified that Bayer decided not to consent to the EPA's direct release of aldicarb toxicology and residue studies to the JMPR. However, the EPA said they would consider if it was possible for them to provide the studies directly to the CCPR without agreement from Bayer.

On September 27, 2019 the EPA stated *"To pursue a definitive decision, CCPR would need to submit a FOIA request, identify the specific studies being requested, and identify all individuals who would have access to the requested studies. EPA would require signed Affirmations of Non-multinational Status from each of those individuals as one of the initial steps after receiving the request. I don't know if EPA would be able to make a final determination on disclosure by December 1, 2019."*

While the studies requested are in the public domain, the only way that the aldicarb studies can be provided to the CCPR is through Freedom of Information requests that are submitted to the EPA by the assigned CCPR reviewers. If this procedure is acceptable Ag Logic Chemical will identify the specific aldicarb studies that should be requested under FOIA. We will include the complete study title and MRID number so that EPA will not have any problem locating the correct studies. We will provide the list of studies to the CCPR so they can submit the FOIA requests.

As it is unlikely that this process can be completed by December 1, 2019, we propose that the aldicarb reevaluation be rescheduled from 2020 to 2021 to allow enough time for the EPA to submit the studies to the CCPR.

Please let us know how the CCPR wishes to proceed and what we can do to ensure the necessary aldicarb studies are available for review.

Best Regards,  
Larry Hodges, Ph.D.  
Director of Regulatory Affairs  
AgLogic Chemical LLC

Phone: 919-932-5800



